

Larry Hogan, Governor · Boyd K. Rutherford, Lt. Governor · Dennis R. Schrader, Secretary

Laboratories Administration Robert A. Myers, Ph.D., Director 1770 Ashland Avenue Baltimore, Maryland 21205

DATE: October 21, 2022

TO: Medical Lab Directors, Local Health Departments, and Healthcare Providers

FROM: Robert A. Myers, Ph.D.

Director, Laboratories Administration

RE: Monkeypox Specimen Collection Updates

The Maryland Department of Health Laboratories Administration is issuing the following Monkeypox specimen collection updates.

See updated specimen collection guidelines. Please note the use of needles or scalpels to deroof the crust/scab is no longer recommended, as per the current CDC guidelines. Blunt-tipped forceps should be used, as needed, to remove the crust/scab of the lesion. The Febrile Rash Submission Form test request is available on the Laboratories Administration's website: https://health.maryland.gov/laboratories/Pages/Home.aspx.

All specimens sent to the Laboratories Administration for orthopox testing still MUST have prior approval for testing from the State Epidemiologists or their authorized designee. Specimens sent without proper approval will not be tested. Please contact the State Epidemiologists at 410-767-6700 (during normal business hours) or 410-795-7365 (after hours) for consultation.

Febrile Rash Low to Moderate Risk Specimen Collection kits are available from the Local Health Departments (LHD's). Please contact your LHD for information regarding specimen collection kits. Please ensure that the kits being used/distributed are within their expiration date. Specimens submitted in expired tubes will be rejected. Unused expired kits are to be returned to the Laboratories Administration for recycling and reuse of the unexpired kit components.

Specimens collected for testing at the Laboratory Administration must be kept refrigerated until delivered to the MD Laboratory within 72 hours of collection. Specimens must be kept frozen (at least -20°C) if they are not received at the MD laboratory within 72 hrs. of collection. This is to ensure the integrity of the specimens received for testing.

For the safety of all who handle specimens submitted for monkey pox testing, specimens must be packaged using Category B shipping requirements, even if specimens are sent via routine or private couriers. See below for basic triple-packaging instructions.

Please contact your Local Health Department or the State Epidemiologists with any other questions or concerns. General questions may be submitted to mdh.monkeypox@maryland.gov.

Guidelines for Orthopox Rule-Out Specimen Submission

MDH Infectious Disease Epidemiology

(Consult Required Prior to specimen submission)

MDH Epi Line (Business Hours) 410-767-6700 MDH Epi/Physician After Hours On-Call at 410-795-7365

MDH Laboratories Administration

Office of Laboratory Emergency Preparedness and Response

Monday – Friday 8:00 a.m. - 4:30 p.m. (Dial in order)

AFTER HOURS (Dial in order)

443-681-3788 - Office phone 410-925-3121 - Cell Phone

410-925-3121 - Cell Phone

443-681-3789 - Office phone

410-408-7521 - Pager

410-408-7521 - Pager

Requested Specimens

Specimen Type	Collection Materials	Instructions	Storage
A minimum of 2* swabs of base of lesion (NO transport media) Required by CDC *Note: One dry swab may be tested at an LRN Reference laboratory for presumptive results. CDC can provide Monkeypox virus-specific testing on the second dry swab specimen if the first dry swab is presumptive positive at the LRN laboratory.	Sterile nylon, polyester, or Dacron swab with a plastic, wood, or thin aluminum shaft. Place in a sterile container with NO transport media. Use 2* swabs per lesion. Place each swab in dry, sterile container and label appropriately. (i.e. body site, description, etc.)	1. Vigorously swab the base of the lesion with a sterile synthetic swab. If scab or crust is removed, collect with forceps. (See below.) If lesion ruptures, use swab to collect lesion fluid. 2. Place swabs in individual sterile containers. DO NOT ADD ANY VIRAL OR UNIVERSAL TRANSPORT MEDIA. 3. Collect specimens from lesions at different anatomic locations if possible. 4. Ensure specimens are appropriately labeled.	Refrigerate after collection at 2-8C. Deliver immediately to MDH.
1 swab of base of a lesion (in viral transport media) Required by MDH (Used for reflex testing if no orthopox viruses found)	Sterile nylon, polyester, or Dacron swab with a plastic, shaft. Tube of viral transport media. Use multiple containers when collecting specimens from multiple lesions.	Same as above EXCEPT place swab in viral transport media in order to allow testing for other viral pathogens if the above specimens test negative for orthopox viruses.	

Specimen Type	Collection Materials	Instructions	Storage
			Conditions
Crust, scab, or biopsy of	Sterile tube with O ring seal	1. If needed, use disposable forceps to	Refrigerate after
lesion		remove the top of the vesicle or pustule	collection at 2-8C.
	(As needed, blunt-tipped	(do not send the forceps). Retain lesion	Deliver
(If able to remove	disposable forceps.)	roof for testing.	immediately to
crust/roof during swab		2. Place specimen in a 1.5 or 2 mL	MDH.
collection, send as	Please note the use of needles or	screw-capped tube with O-ring. DO	
additional specimen)	scalpels to de-roof the crust/scab	NOT ADD ANY VIRAL TRANSPORT	Specimens may be
	is no longer recommended, as per	MEDIA.	kept at 2-8C for up
	the current CDC guidelines.	3. Collect specimens from lesions at	to 72 hrs, after
		different anatomic locations if possible.	which point they
		4. Ensure specimen are appropriately	must be frozen at
		labeled.	minimum -20C.
			They may stay
			frozen for up to
			two months.

Required Personal Protective Equipment (PPE) to collect specimens

- Disposable Gown
- Gloves
- Fit tested N95 or PAPR *Surgical mask if not available
- Eye protection- face shield or goggles (note that eye-glasses are not eye protection)

Alcohol based hand sanitizer or sink with soap and water for hand hygiene

Required Data Elements on Specimens and Requisition Forms

- All specimens may be listed out on the FEBRILE RASH (LOW TO MODERATE RISK) form (Appendix 2). Use one Tube ID for each body site/specimen type.
- Each individual specimen must be labeled with the following:
 - o Patient name
 - o DOB
 - Collection date
 - o Specimen site
- Unlabeled specimens will be AUTOMATICALLY REJECTED.
- Under Presumptive Clinical Diagnosis, specify Other: Monkeypox
- Test Request Authorized By is your ordering provider. Please only list an MD, DO, PA-C, or CRNP.
- Health Care Provider is the submitting facility name.
- All patient demographic elements are required.
- Consult with MDH Epidemiology to determine risk category.
- Ship packages using Category B Shipping (See below).
- Address package to MDH Labs, "For Monkey Pox testing, ATTN:Amy Armitage or Sali Lukula"

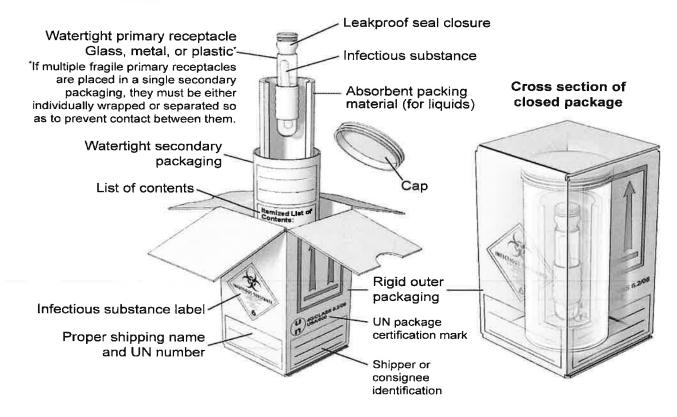
Appendix 1: Packaging and Shipping

Specimens must be shipped using Category B shipping requirements.

Refer to the attached document (Basic Triple Packaging) for packaging and shipping guidelines. If necessary, the MDH Laboratories Administration will arrange for an emergency courier, but will only do so after consultation with the patient's physician and MDH Physician-On-Call or Epidemiologist.

BASICTRIPLE PACKAGING

Basic triple packaging systems include a primary receptacle such as a tube with adhesive tape around the screw cap or a plate with parafilm around the edges. The primary (1°) receptacle, along with required absorbent and cushioning material, is placed inside a secondary (2°) container. The 2° container for diagnostic specimens should be a sealed biohazard or Ziploc bag. The 2° container is then securely placed within an outer shipping container (tertiary (3°) container), generally a corrugated cardboard box with cushioning material inside to surround the 2° container. This outermost container bears the name, address, and telephone number of shipper, name of person responsible with 24/7 telephone number, and the complete name, shipping address, and telephone number of the recipient, plus all the required markings. Include an itemized list of contents in a sealed plastic bag, placed between the 3° and 2° containers.



BASICTRIPLE PACKAGING:

- A watertight primary receptacle.
- A watertight secondary receptacle.
- An outer packaging of adequate strength for its capacity, mass and intended use.

Note: For a liquid specimen, absorbent material must be placed between the primary and secondary containers and be capable of absorbing the entire contents of the primary receptacle(s).

Certified packaging systems are designed to withstand specific pressure changes and drop tests. Packaging systems that meet the

packing instruction standards are currently available from vendors specializing in products certified to meet the IATA, USPS, and other carriers' requirements. Packaging systems using fiberboard or aluminum canisters, zip-lock bags, or other uncertified components may not be in compliance.

IT IS THE RESPONSIBILITY OF THE SHIPPER TO COMPLY WITH ALL LAWS AND REGULATIONS REGARDING THE SHIPPING OF INFECTIOUS SUBSTANCES.



Laboratories Administration 1770 Ashland Ave Baltimore, Maryland 21205

Robert A. Myers, Ph.D., Director

FEBRILE RASH (Low and Moderate Risk)

		Health Care Provider					
		Address					
		City	County				
		State	Zip Code				
		Contact Name					
		Phone#	Fax#				
		Test Request Authorized by:					
		_	TYPE OR PRINT				
	Patient's SS# (last 4 digits) Case# Patient Lab No Last Name First Middle						
		of Birth//	Sex M F				
	Addre	ess					
	City_	Co	ountyState	Zip			
		Risk Category:	Low Risk Moderate Risk				
	For		Risk testing, collect the following s ACE ONLY ONE LESION PER TUE	=			
ube id	collection device	specimen type needed	body site of collection (arm, chest, face, etc.)	description of site (vesicle, pustule, etc.)			
1	tube with	swab of base of lesion					
2	transport media	swab of base of lesion					
3	long tube with swab	swab of base of lesion					
4	and no liquid	swab of base of lesion					
5		crusUscab of lesion					
6	small empty tube	crusUscab of lesion					
7	with O ring seal	crusUscab of lesion					
8		crusUscab of lesion					
	Presumptive	Clinical Diagnosis: Ch	ickenpox Herpesvirus S	malipox			
	Smallpox vaccine (Vaccinia) Other:(specify)						
	Date of Onse	t:// (Month / Day / Year)	- 2				
	Date Specime	en Collected	Reported				

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